IN THE CLAIMS

1-27. (canceled)

- 28. (currently amended) A composition comprising a cell in which a molecular complex is bound to the surface of the cell, wherein the molecular complex comprises at least two first fusion proteins and at least two second four fusion proteins, wherein:
- (a) each of the two first fusion proteins comprises comprises an immunoglobulin heavy chain, wherein the immunoglobulin heavy chain comprises a variable region, and an extracellular portion of a first transmembrane polypeptide; and
- (b) <u>each of the</u> two second fusion proteins <u>comprises</u> an immunoglobulin light chain and an extracellular portion of a second transmembrane polypeptide;

wherein the <u>at least two first fusion proteins</u> and the at least two second fusion proteins associate to form a <u>the</u> molecular complex, wherein the molecular complex comprises two ligand binding sites, <u>wherein</u> each ligand binding site <u>is</u> formed by the extracellular <u>domain</u> domains of a <u>the</u> first <u>transmembrane polypeptide</u> and <u>the extracellular domain of a second transmembrane polypeptide</u> polypeptides, wherein the affinity of the molecular complex for a cognate ligand is increased at least two-fold over a dimeric molecular complex consisting of a <u>the</u> first and a <u>the</u> second fusion protein.

- 29. (original) The composition of claim 28 wherein the first transmembrane polypeptide is an MHC class IIβ chain and wherein the second transmembrane polypeptide is an MHC class IIα chain.
- 30. (original) The composition of claim 28 wherein the first transmembrane polypeptide is a TCR α chain and wherein the second transmembrane polypeptide is a TCR β chain.

- 31. (original) The composition of claim 28 further comprising a pharmaceutically acceptable carrier.
- 32. (previously amended) The composition of claim 28 wherein a population of the molecular complexes is bound to the cell, wherein an identical antigenic peptide is bound to each ligand binding site.
 - 33-50. (canceled)
- 51. (previously added) The composition of claim 28 wherein the immunoglobulin heavy chain is an IgG1 heavy chain.
- 52. (previously added) The composition of claim 28 wherein the immunoglobulin light chain is an Igk chain.
- 53. (previously added) The composition of claim 28 wherein the first fusion proteins comprise a first peptide linker between the immunoglobulin heavy chain and the extracellular domain of the first transmembrane polypeptide and wherein the second fusion proteins comprise a second peptide linker between the immunoglobulin light chain and the extracellular domain of the second transmembrane polypeptide.
- 54. (previously added) The composition of claim 53 wherein the first peptide linker is GLY-GLY-THR-SER-GLY (SEQ ID NO:10).
- 55. (previously added) The composition of claim 53 wherein the second peptide linker is GLY-SER-LEU-GLY-SER (SEQ ID NO:11).
- 56. (previously amended) The composition of claim 32 wherein the antigenic peptides are bound to the ligand binding sites by a method comprising the step of:

incubating the cell in the presence of the antigenic peptides, whereby the antigenic peptides are bound to the ligand binding sites.

- 57. (previously amended) The composition of claim 32 wherein the antigenic peptides are bound to the ligand binding sites by a method comprising the steps of:
 - (a) alkaline stripping of the molecular complex to provide an alkaline stripped molecular complex;
 - (b) neutralization of the alkaline stripped molecular complex to provide a neutralized molecular complex;
 - (c) incubation of the neutralized molecular complex in the presence of an excess of the antigenic peptides; and
 - (c) slow refolding of the neutralized molecular complex in the presence of the excess of the antigenic peptides.
- 58. (previously added) The composition of claim 32 wherein the antigenic peptides are covalently bound.
- 59. (previously added) The composition of claim 28 wherein the molecular complex is conjugated to a toxin.
- 60. (previously amended) The composition of claim 28 wherein the molecular complex is conjugated to a lymphokine or other effector molecule which stimulates an immune response.)